WHAT IS CLAIMED IS:

1. A method for treating an inflammatory disease or reducing an inflammatory reaction, said method comprising: administering a SOC inhibitor, thereby treating said inflammatory disease or reducing said inflammatory reaction.

2. The method of claim 1, wherein said inflammatory disease or inflammatory reaction is a skin disorder.

3. The method of claim 2, wherein said skin disorder is selected from the group consisting of atopic dermatitis, psoriasis, neurogenic inflammation, skin photodamage, a cell carcinoma, keratosis, and a disorder of keratinization.

4. The method of claim 1, wherein said inflammatory disease or inflammatory reaction is an inflammatory pulmonary disease or reaction.

5. The method of claim 4, wherein said inflammatory pulmonary disease or reaction is selected from the group consisting of asthma, allergic rhinitis, chronic obstructive pulmonary disease and adult respiratory distress syndrome.

6. The method of claim 1, wherein said inflammatory disease or inflammatory reaction is an inflammatory musculoskeletal disease or reaction.

7. The method of claim 6, wherein said inflammatory musculoskeletal disease is a member selected from the group consisting of psoriatic arthritis, osteoarthritis, and osteoporosis.

8. The method of claim 1, wherein said inflammatory disease or inflammatory reaction is an inflammatory gastrointestinal or urogenital disease or reaction.

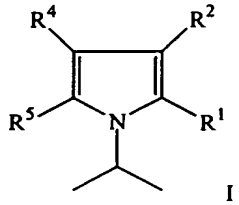
9. The method of claim 8, wherein said inflammatory gastrointestinal or urogenital disease or reaction is a member selected from the group consisting of inflammatory bowel disease, enterocolitis, gastritis, vaginitis, and interstitial cystitis.

10. The method of claim 1, wherein said inflammatory disease or inflammatory reaction is an autoimmune disease or reaction.

11. The method of claim 10, wherein said autoimmune disease is a member selected from the group consisting of multiple sclerosis, type II diabetes, lupus, and rheumatoid arthritis.

12. The method of claim 1, wherein said inflammatory disease or inflammatory reaction is transplantation treatment.

13. The method of claim 1, wherein said SOC inhibitor is a compound having the formula:



I

wherein:

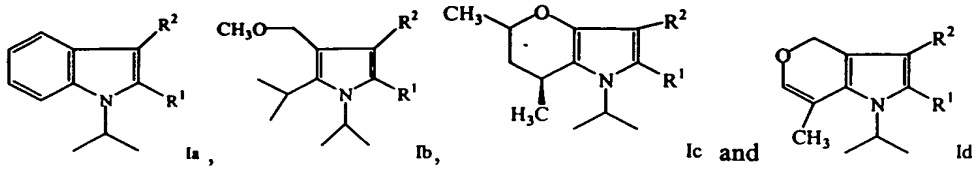
R^1 is a member selected from the group consisting of optionally substituted alkyl, optionally substituted alkoxy, optionally substituted alkylene having at least 2 sites of unsaturation, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, optionally substituted heteroarylthioalkyl, optionally substituted heteroaryliminooxyalkyl, optionally substituted heterocyclyl, optionally substituted oximinoaryl and optionally substituted heteroarylalkoxy;

R^2 is a member selected from the group consisting of optionally substituted $(C_1\text{-}C_6)$alkyl, optionally substituted $(C_1\text{-}C_6)$alkoxy, acyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy;

R^4 is an optionally substituted alkyl; and

R^5 is an optionally substituted alkyl, or alternatively, R^4 and R^5 and the carbons to which they are attached, joined to form an optionally substituted aryl or optionally substituted heteroalkyl 5-or 6 membered ring.

14. The method of claim 13, wherein said compound is a member selected from the group consisting of



Ia, Ib, Ic and Id

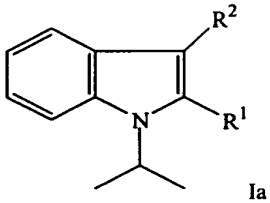
wherein:

R^1 is a member selected from the group consisting of optionally substituted alkyl, optionally substituted alkoxy, optionally substituted (C_2-C_{18})alkylene having at least 2 sites of unsaturation, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, optionally substituted heteroarylalkoxy, and optionally substituted; and

R^2 is a member selected from the group consisting of optionally substituted (C_1-C_6)alkyl, optionally substituted (C_1-C_6)alkoxy, acyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy.

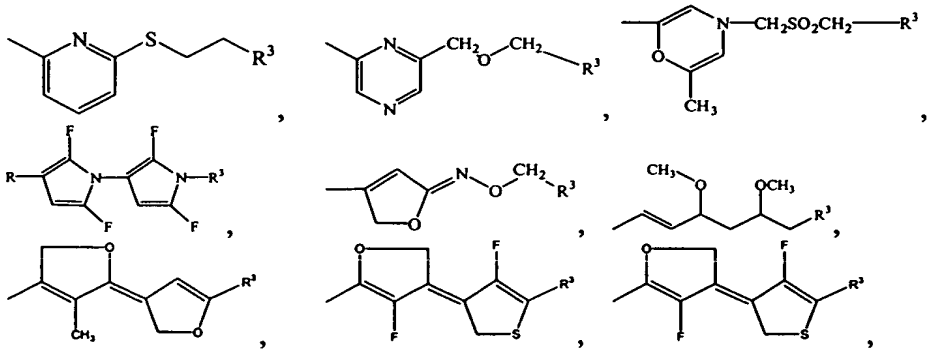
15. The method of claim **13**, wherein R^1 is an optionally substituted (C_2-C_{18})alkylene having at least 2 sites of unsaturation.

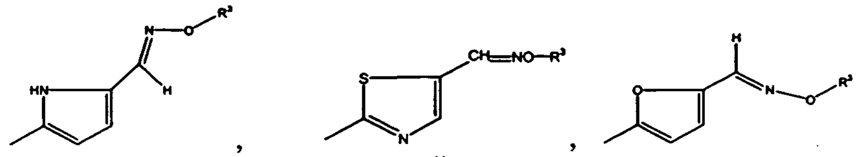
16. The method of claim **14**, wherein said compound is

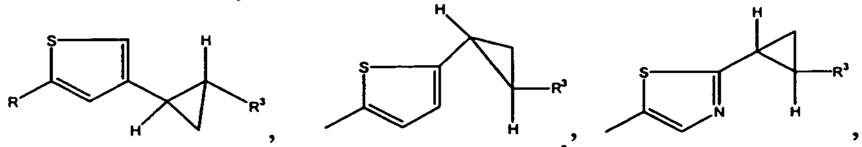


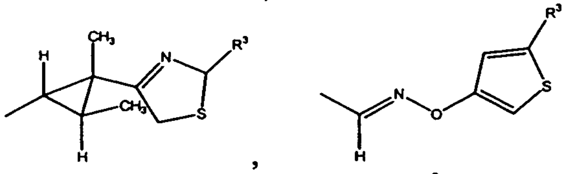
Ia

17. The method of claim **13**, wherein R^1 is a member selected from the group consisting of



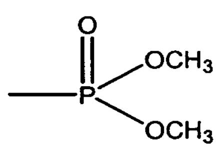
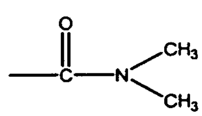
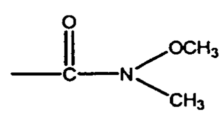
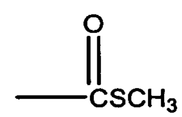


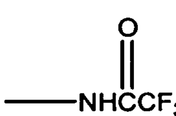
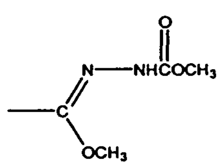
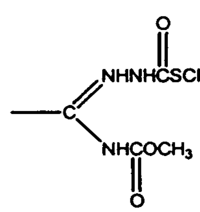
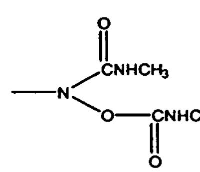


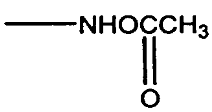
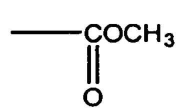
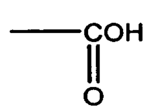
 , $—(CH{=}CH)_2—(CH_2)_x—R^3$ wherein x is about 1 to about 14 and $—(CH{=}CH)_3—R^3$; and

R^3 is a member selected from the group consisting of consisting of alkyl, alkoxysulfonyl, dialkylphosphono, optionally substituted carbamoyl, alkylthiocarbonyl, optionally substituted alkylamido, optionally substituted amidino, optionally substituted alkylsulfonamido, alkoyloxyamino, alkylaminosulfonamido, and alkoxycarbonyl.

18. The method of claim **17**, wherein R^3 is a member selected from the group consisting of consisting of

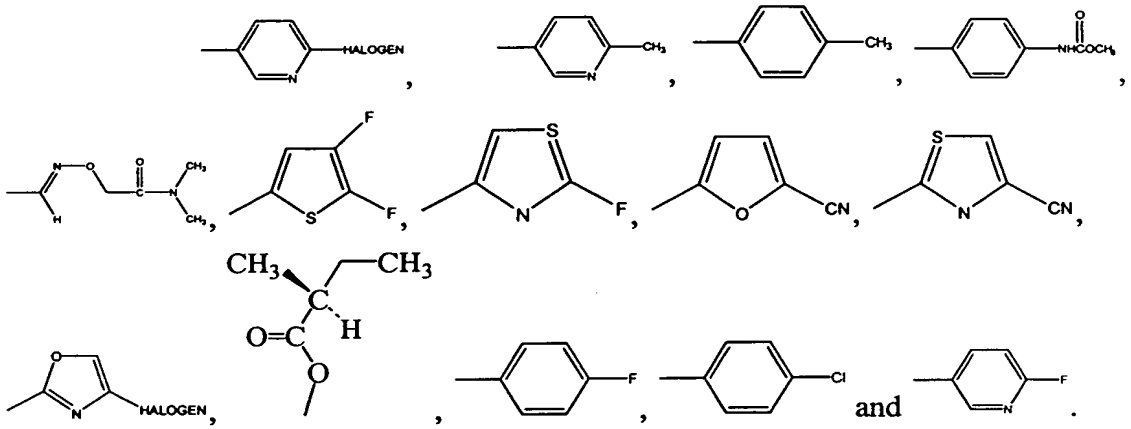
$—SO_2OCH_3$,  ,  ,  ,  ,

 ,  ,  , $—NHSO_2CF_3$,  ,

 , $—NHSO_2NHCH_3$,  ,  , $-(CH_2)_nCH_3$ wherein n is about 1 to about 10 and $-CH_3$.

19. The method of claim **18**, wherein R^2 is independently a member selected from the group consisting of optionally substituted alkoxy, acyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy.

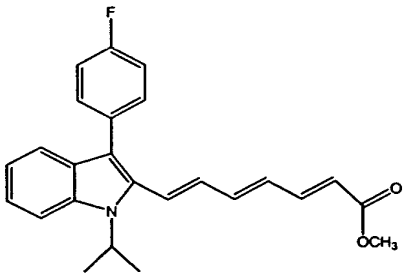
20. The method of claim **19,** wherein R^2 is a member selected from the group consisting of consisting of



and

21. The method of claim **20,** wherein R^2 is p-fluorophenyl.

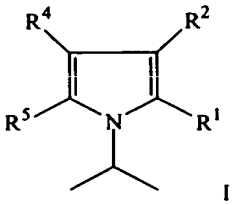
22. The method of claim **13,** wherein said compound has the formula



23. The method of claim **1,** wherein said SOC inhibitor is a δ-lactone-containing statin.

24. The method of claim **23,** wherein said δ-lactone-containing statin is a member selected from the group consisting of lovastatin, mevastatin, and simvastatin.

25. A compound having the formula



I

wherein:

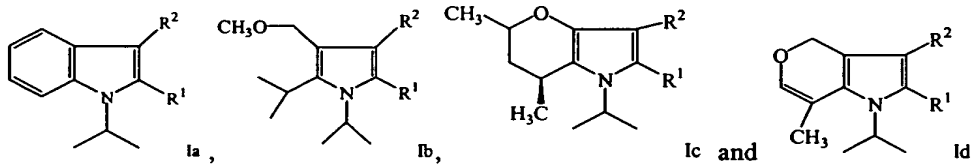
R^1 is a member selected from the group consisting of optionally substituted alkyl, optionally substituted alkoxy, optionally substituted alkylene having at least 2 sites of unsaturation, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, optionally substituted heteroarylthioalkyl, optionally substituted heteroaryliminooxyalkyl, optionally substituted heterocyclyl, optionally substituted oximinoaryl and optionally substituted heteroarylalkoxy;

R^2 is a member selected from the group consisting of optionally substituted $(C_1\text{-}C_6)$alkyl, optionally substituted $(C_1\text{-}C_6)$alkoxy, acyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy;

R^4 is an optionally substituted alkyl; and

R^5 is an optionally substituted alkyl, or alternatively, R^4 and R^5 and the carbons to which they are attached, joined to form an optionally substituted aryl or optionally substituted heteroalkyl 5-or 6 membered ring.

26. The compound of claim **25,** wherein said compound is a member selected from the group consisting of



Ia, Ib, Ic and Id

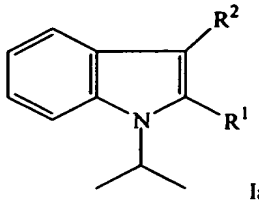
wherein:

R^1 is a member selected from the group consisting of optionally substituted alkyl, optionally substituted alkoxy, optionally substituted $(C_2\text{-}C_{18})$alkylene having at least 2 sites of unsaturation, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, optionally substituted heteroarylalkoxy, and optionally substituted; and

R^2 is a member selected from the group consisting of optionally substituted $(C_1\text{-}C_6)$alkyl, optionally substituted $(C_1\text{-}C_6)$alkoxy, acyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy.

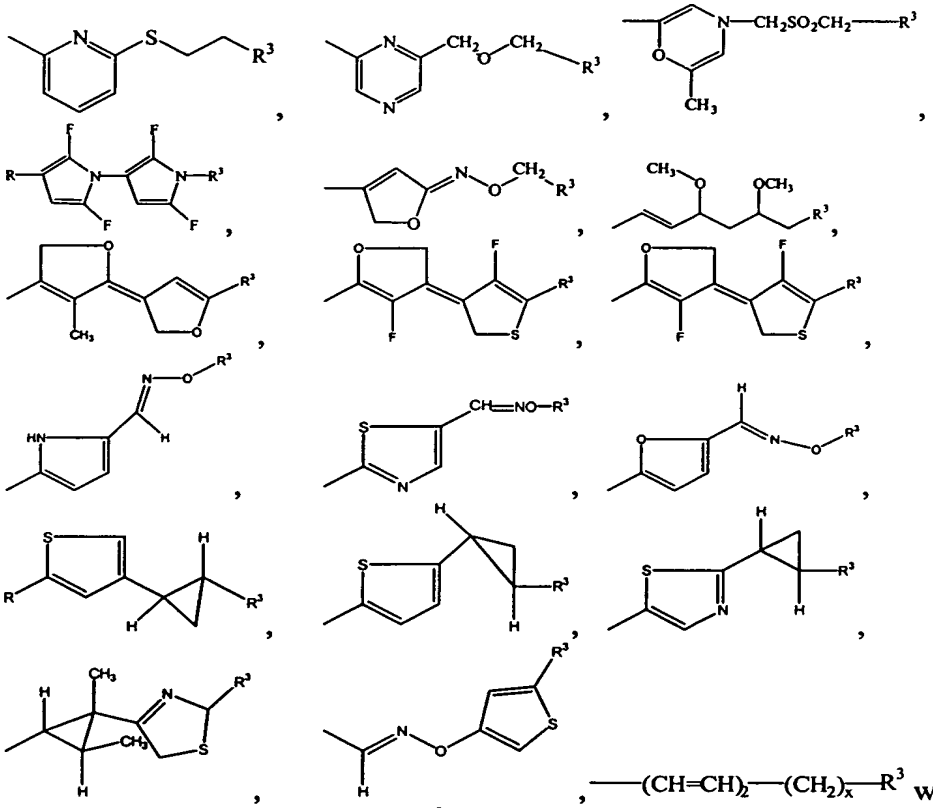
27. The compound of claim **25,** wherein R^1 is an optionally substituted $(C_2\text{-}C_{18})$alkylene having at least 2 sites of unsaturation.

28. The compound of claim **26**, wherein said compound is



Ia

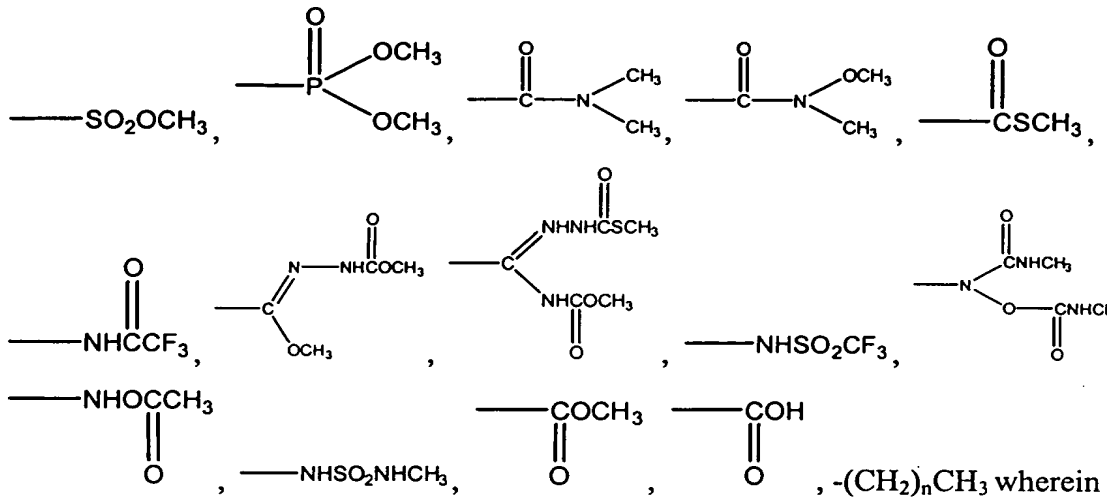
29. The compound of claim **25**, wherein R^1 is a member selected from the group consisting of



wherein x is about 1 to about 14 and $-(CH=CH)_3-R^3$; and

R^3 is a member selected from the group consisting of consisting of alkyl, alkoxysulfonyl, dialkylphosphono, optionally substituted carbamoyl, alkylthiocarbonyl, optionally substituted alkylamido, optionally substituted amidino, optionally substituted alkylsulfonamido, alkoyloxyamino, alkylaminosulfonamido, and alkoxycarbonyl.

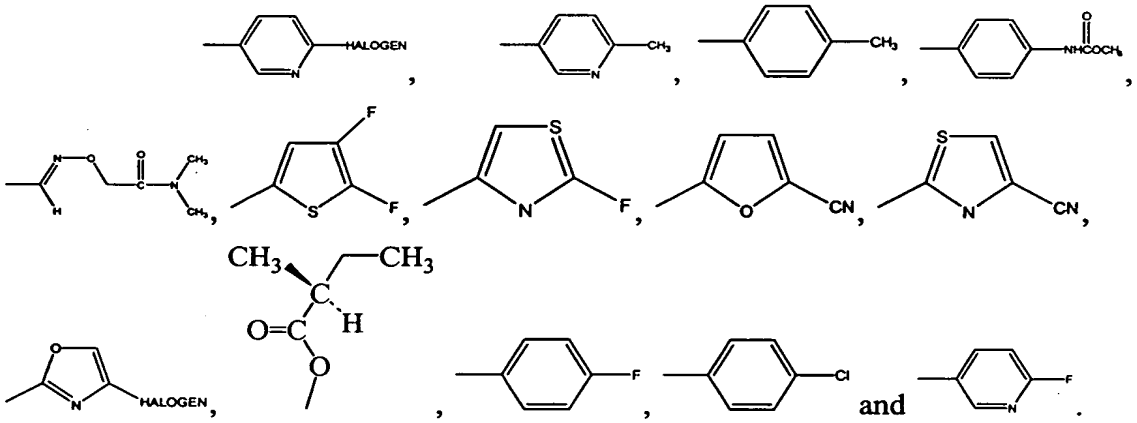
30. The compound of claim **29**, wherein R^3 is a member selected from the group consisting of consisting of



, $-(CH_2)_nCH_3$ wherein n is about 1 to about 10 and $-CH_3$.

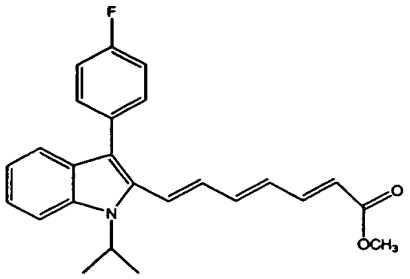
31. The compound of claim **29**, wherein R^2 is independently a member selected from the group consisting of optionally substituted alkoxy, acyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy.

32. The compound of claim **31**, wherein R^2 is a member selected from the group consisting of consisting of

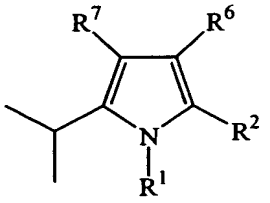


33. The compound of claim **32**, wherein R^2 is p-fluorophenyl.

34. The compound of claim **25**, wherein said compound has the formula



35. A compound having the formula



II

wherein:

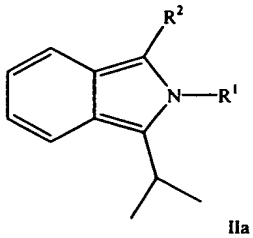
R^1 is a member selected from the group consisting of optionally substituted alkyl, optionally substituted alkoxy, optionally substituted alkylene having at least 2 sites of unsaturation, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, optionally substituted heteroarylthioalkyl, optionally substituted heteroaryliminooxyalkyl, optionally substituted heterocyclyl, optionally substituted oximinoaryl and optionally substituted heteroarylalkoxy;

R^2 is independently a member selected from the group consisting of optionally substituted $(C_1\text{-}C_6)$alkyl, optionally substituted $(C_1\text{-}C_6)$alkoxy, acyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy;

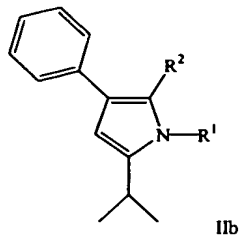
R^6 is a member selected from the group consisting of an optionally substituted alkyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy; and

R^7 is a member selected from the group consisting of an optionally substituted alkyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy; or alternatively, R^6 and R^7 and the carbons to which they are attached, joined to form an optionally substituted aryl or optionally substituted heteroalkyl 5-or 6 membered ring.

36. The compound of claim 35, wherein said compound is a member selected from the group consisting of


IIa

and

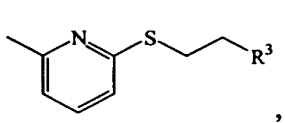

IIb

wherein:

R^1 is a member selected from the group consisting of optionally substituted alkyl, optionally substituted alkoxy, optionally substituted (C_2-C_{18})alkylene having at least 2 sites of unsaturation, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy; and

R^2 is independently a member selected from the group consisting of optionally substituted (C_1-C_6)alkyl, optionally substituted (C_1-C_6)alkoxy, acyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy.

37. The compound of claim 36, wherein R^1 is a member selected from the group consisting of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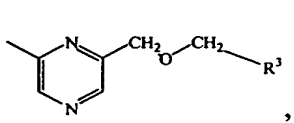
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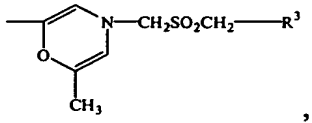
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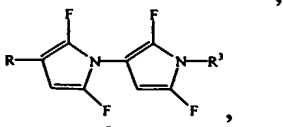
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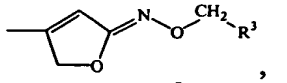
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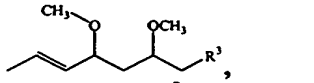
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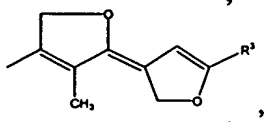
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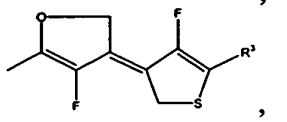
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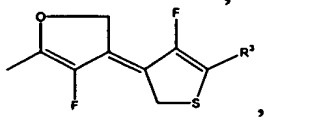
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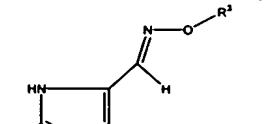
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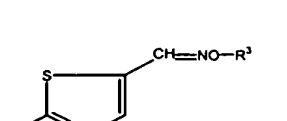
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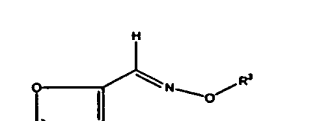
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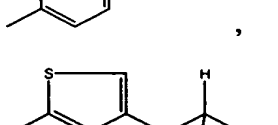
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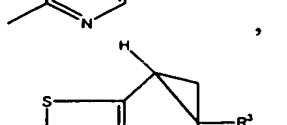
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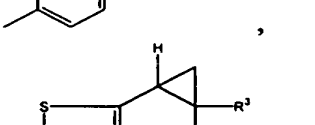
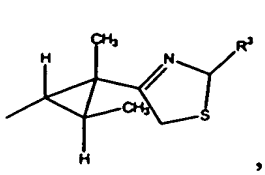
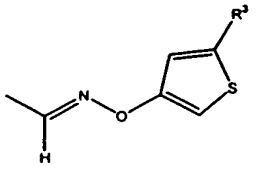
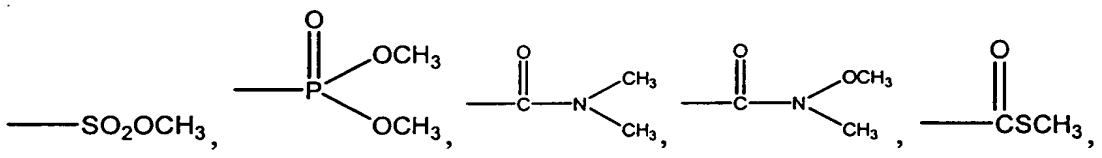
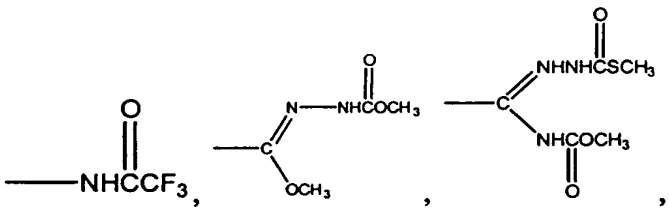
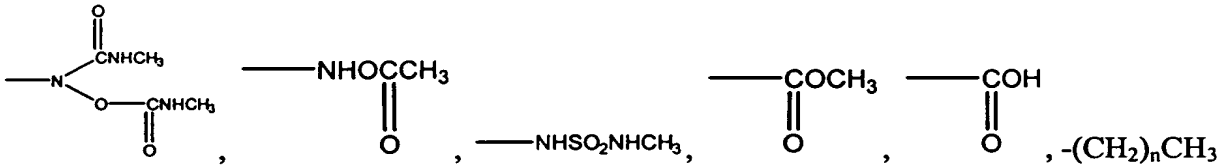

,

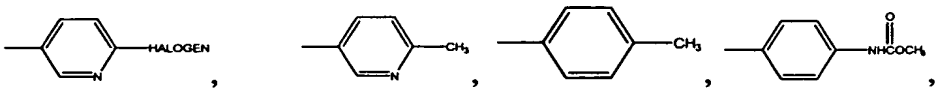
about 1 to about 14 and ; and

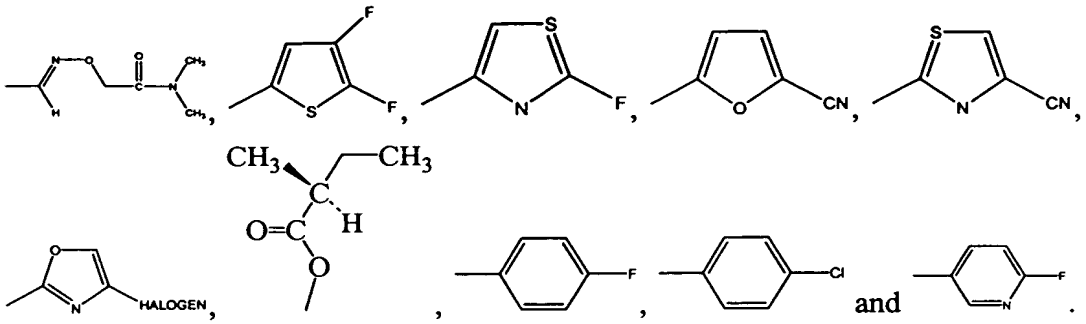
R^3 is a member selected from the group consisting of consisting of alkyl, alkoxysulfonyl, dialkylphosphono, optionally substituted carbamoyl, alkylthiocarbonyl, optionally substituted alkylamido, optionally substituted amidino, optionally substituted alkylsulfonamido, alkoyloxyamino, alkylaminosulfonamido, and alkoxycarbonyl.

38. The compound of claim **37,** wherein R^3 is a member selected from the group consisting of consisting of




wherein n is about 1 to about 10 and $-CH_3$.

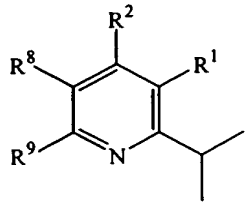
39. The compound of claim **36,** wherein R^2 is a member selected from the group consisting of consisting of





40. The compound of claim 39, wherein R^2 is p-fluorophenyl.

41. A compound having the formula



III

wherein:

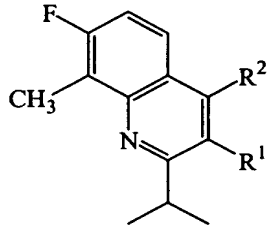
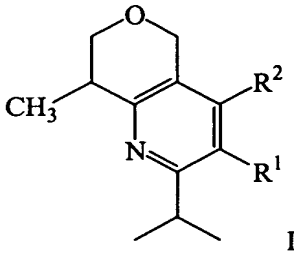
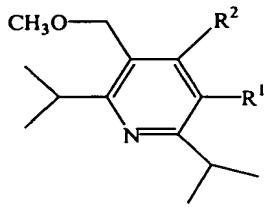
R^1 is a member selected from the group consisting of optionally substituted alkyl, optionally substituted alkoxy, optionally substituted alkylene having at least 2 sites of unsaturation, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, optionally substituted heteroarylthioalkyl, optionally substituted heteroaryliminooxyalkyl, optionally substituted heterocyclyl, optionally substituted oximinoaryl and optionally substituted heteroarylalkoxy;

R^2 is independently a member selected from the group consisting of optionally substituted (C_1-C_6)alkyl, optionally substituted (C_1-C_6)alkoxy, acyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy;

R^8 is a member selected from the group consisting of an optionally substituted alkyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy;

R^9 is a member selected from the group consisting of an optionally substituted alkyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy; or alternatively, R^8 and R^9 and the carbons to which they are attached, joined to form an optionally substituted aryl or optionally substituted heteroalkyl 5-or 6 membered ring.

42. The compound of claim **41**, wherein said compound is a member selected from the group consisting of

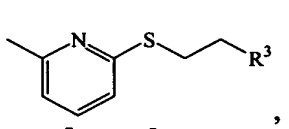

IIIa,

IIIb, and

IIIc

wherein:

R^1 is a member selected from the group consisting of optionally substituted alkyl, optionally substituted alkoxy, optionally substituted (C_2-C_{18})alkylene having at least 2 sites of unsaturation, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, optionally substituted heteroarylalkoxy, and optionally substituted; and

R^2 is independently a member selected from the group consisting of optionally substituted (C_1-C_6)alkyl, optionally substituted (C_1-C_6)alkoxy, acyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy.

43. The compound of claim **42**, wherein R^1 is a member selected from the group consisting of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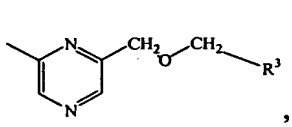
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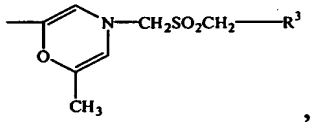
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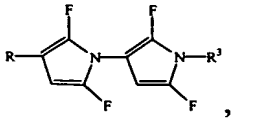
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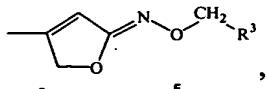
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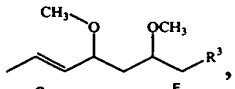
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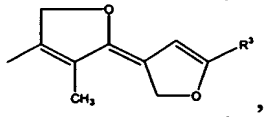
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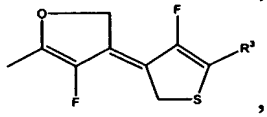
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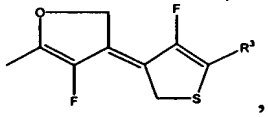
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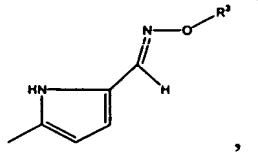
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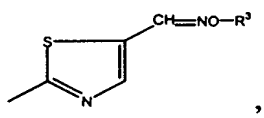
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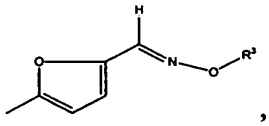
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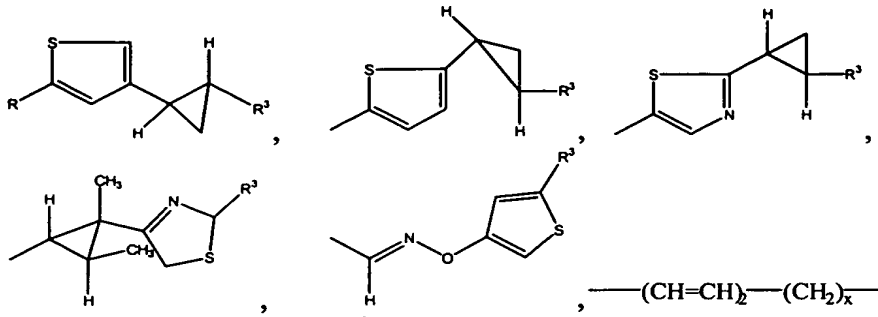
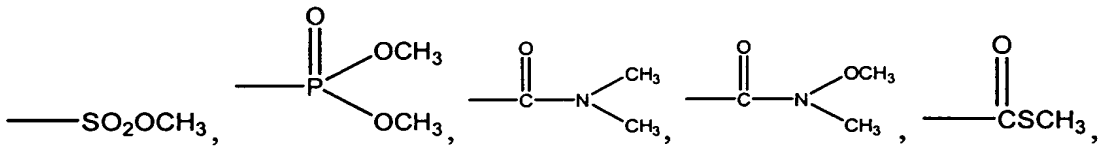
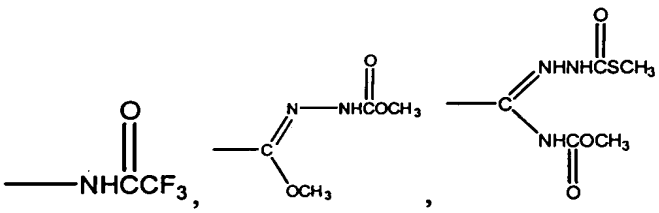


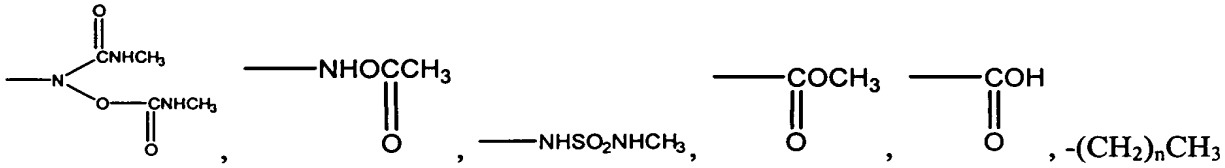
, , , , , —(CH=CH)$_2$—(CH$_2$)$_x$—R^3 wherein x is about 1 to about 14 and —(CH=CH)$_3$—R^3; and

R^3 is a member selected from the group consisting of consisting of alkyl, alkoxysulfonyl, dialkylphosphono, optionally substituted carbamoyl, alkylthiocarbonyl, optionally substituted alkylamido, optionally substituted amidino, optionally substituted alkylsulfonamido, alkoyloxyamino, alkylaminosulfonamido, and alkoxycarbonyl.

44. The compound of claim **43,** wherein R^3 is a member selected from the group consisting of consisting of

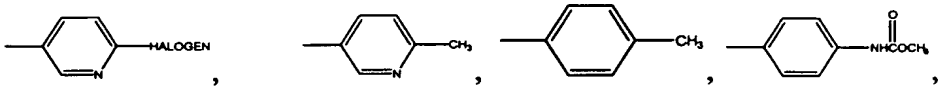


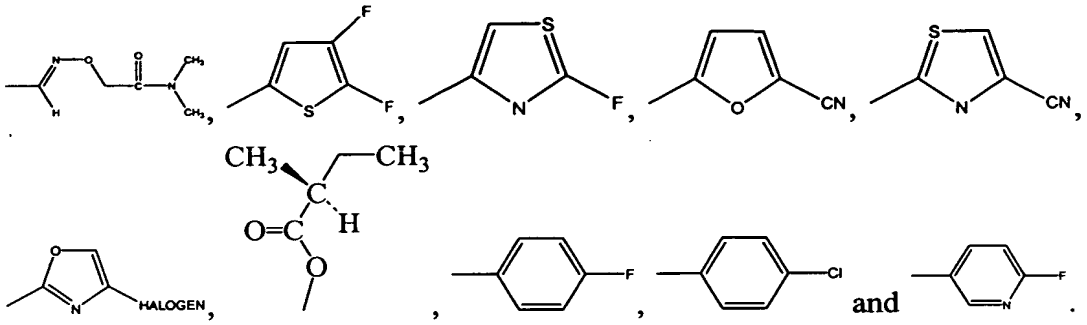




wherein n is about 1 to about 10 and -CH$_3$.

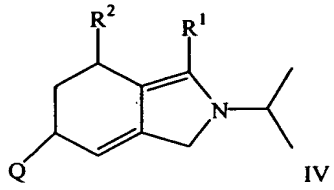
45. The compound of claim **42,** wherein R^2 is a member selected from the group consisting of consisting of





46. The compound of claim **45**, wherein R^2 is p-fluorophenyl.

47. A compound having the formula



IV

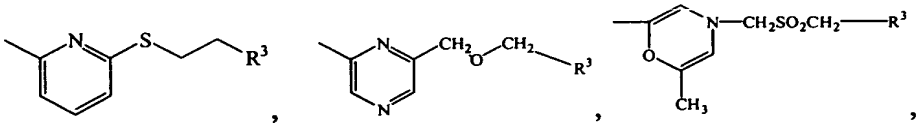
wherein:

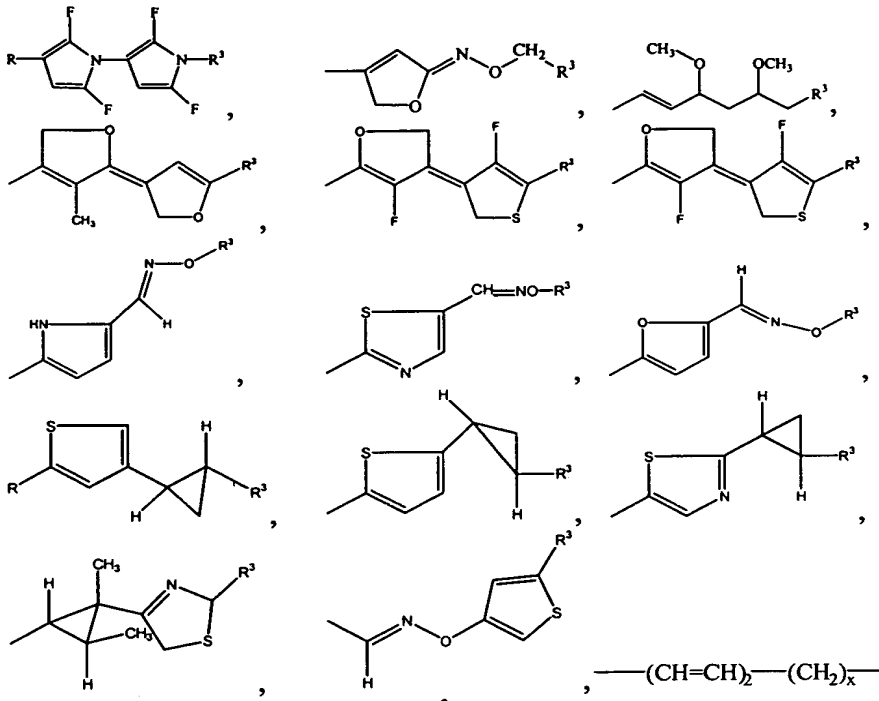
R^1 is a member selected from the group consisting of optionally substituted alkyl, optionally substituted alkoxy, optionally substituted alkylene having at least 2 sites of unsaturation, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, optionally substituted heteroarylthioalkyl, optionally substituted heteroaryliminooxyalkyl, optionally substituted heterocyclyl, optionally substituted oximinoaryl and optionally substituted heteroarylalkoxy;

R^2 is a member selected from the group consisting of optionally substituted $(C_1\text{-}C_6)$alkyl, optionally substituted $(C_1\text{-}C_6)$alkoxy, acyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy; and

Q is a member selected from the group consisting of hydrogen, optionally substituted alkyl, optionally substituted alkoxy and hydroxy.

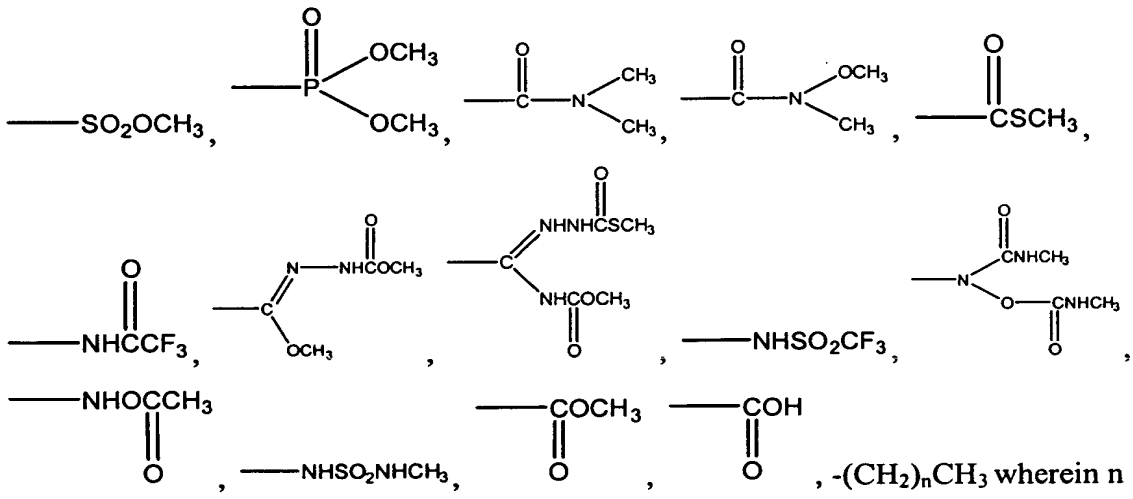
48. The compound of claim **47**, wherein R^1 is a member selected from the group consisting of



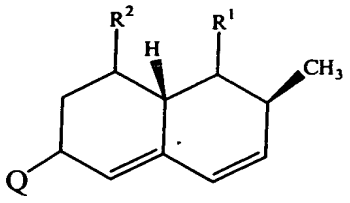

, —(CH=CH)$_2$—(CH$_2$)$_x$—R^3 wherein x is about 1 to about 14 and —(CH=CH)$_3$—R^3; and

R^3 is a member selected from the group consisting of consisting of alkyl, alkoxysulfonyl, dialkylphosphono, optionally substituted carbamoyl, alkylthiocarbonyl, optionally substituted alkylamido, optionally substituted amidino, optionally substituted alkylsulfonamido, alkoyloxyamino, alkylaminosulfonamido, and alkoxycarbonyl.

49. The compound of claim **48,** wherein R^3 is a member selected from the group consisting of consisting of


, -(CH$_2$)$_n$CH$_3$ wherein n is about 1 to about 10 and -CH$_3$.

50. A compound having the formula



V

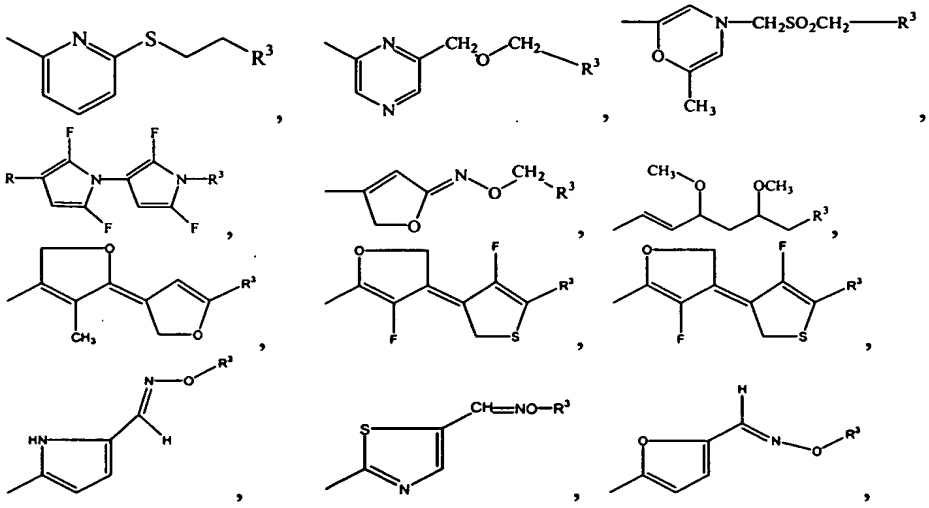
wherein:

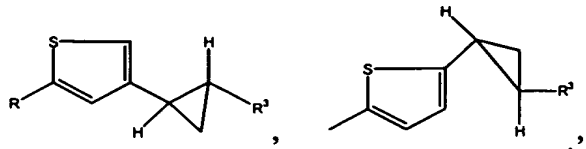
R^1 is a member selected from the group consisting of optionally substituted alkyl, optionally substituted alkoxy, optionally substituted alkylene having at least 2 sites of unsaturation, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, optionally substituted heteroarylthioalkyl, optionally substituted heteroaryliminooxyalkyl, optionally substituted heterocyclyl, optionally substituted oximinoaryl and optionally substituted heteroarylalkoxy;

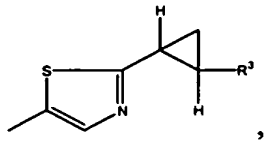
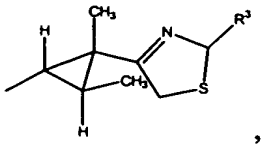
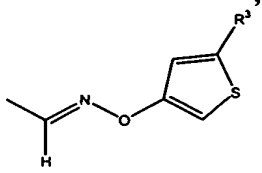
R^2 is independently a member selected from the group consisting of optionally substituted $(C_1\text{-}C_6)$alkyl, optionally substituted $(C_1\text{-}C_6)$alkoxy, acyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy; and

Q is a member selected from the group consisting of hydrogen, optionally substituted alkyl, optionally substituted alkoxy and hydroxy.

51. The compound of claim **50,** wherein R^1 is a member selected from the group consisting of

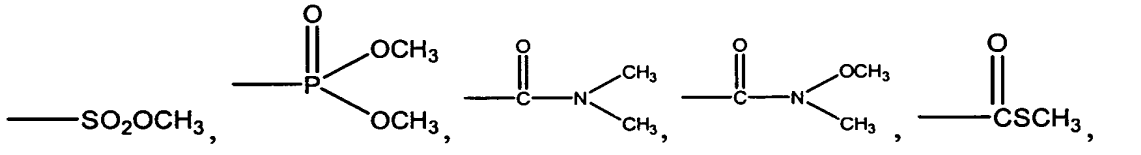
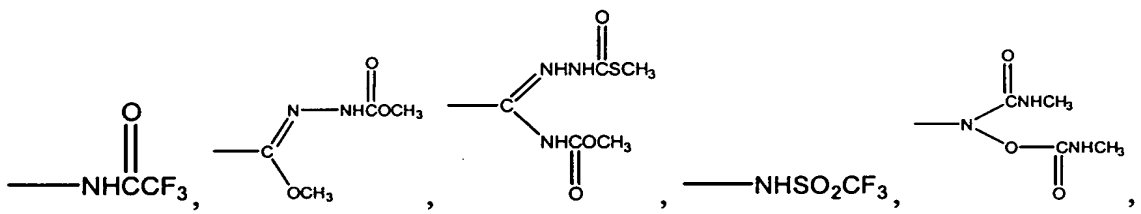
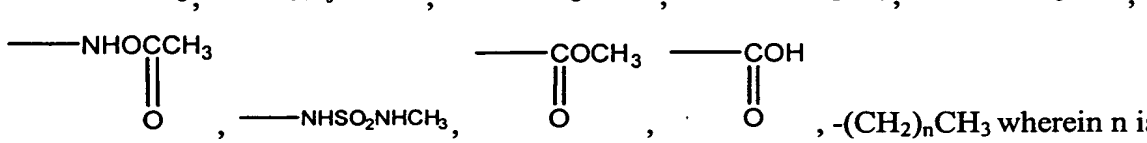







and —(CH=CH)$_3$—R^3; and

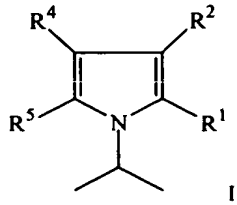
R^3 is a member selected from the group consisting of consisting of alkyl, alkoxysulfonyl, dialkylphosphono, optionally substituted carbamoyl, alkylthiocarbonyl, optionally substituted alkylamido, optionally substituted amidino, optionally substituted alkylsulfonamido, alkoyloxyamino, alkylaminosulfonamido, and alkoxycarbonyl.

52. The compound of claim **51**, wherein R^3 is a member selected from the group consisting of consisting of




, -(CH$_2$)$_n$CH$_3$ wherein n is about 1 to about 10 and -CH$_3$.

53. A pharmaceutical composition, said pharmaceutical composition comprising:

a compound having the formula


I

wherein:

R^1 is a member selected from the group consisting of optionally substituted alkyl, optionally substituted alkoxy, optionally substituted alkylene having at least 2 sites of unsaturation, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, optionally substituted heteroarylthioalkyl, optionally substituted heteroaryliminooxyalkyl, optionally substituted heterocyclyl, optionally substituted oximinoaryl and optionally substituted heteroarylalkoxy;

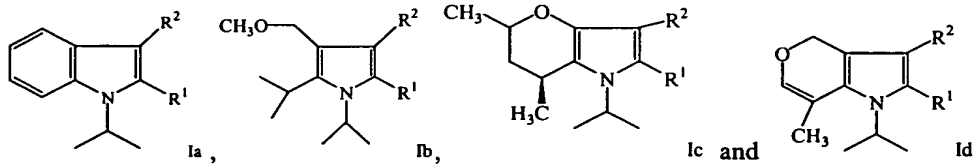
R^2 is a member selected from the group consisting of optionally substituted $(C_1\text{-}C_6)$alkyl, optionally substituted $(C_1\text{-}C_6)$alkoxy, acyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy;

R^4 is an optionally substituted alkyl;

R^5 is an optionally substituted alkyl, or alternatively, R^4 and R^5 and the carbons to which they are attached, joined to form an optionally substituted aryl or optionally substituted heteroalkyl 5-or 6 membered ring; and

a pharmaceutically acceptable excipient therefor.

54. The pharmaceutical composition of claim **53**, wherein said compound is a member selected from the group consisting of



Ia, Ib, Ic and Id

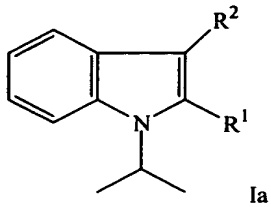
wherein:

R^1 is a member selected from the group consisting of optionally substituted alkyl, optionally substituted alkoxy, optionally substituted $(C_2\text{-}C_{18})$alkylene having at least 2 sites of unsaturation, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, optionally substituted heteroarylalkoxy, and optionally substituted; and

R^2 is independently a member selected from the group consisting of optionally substituted $(C_1\text{-}C_6)$alkyl, optionally substituted $(C_1\text{-}C_6)$alkoxy, acyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy.

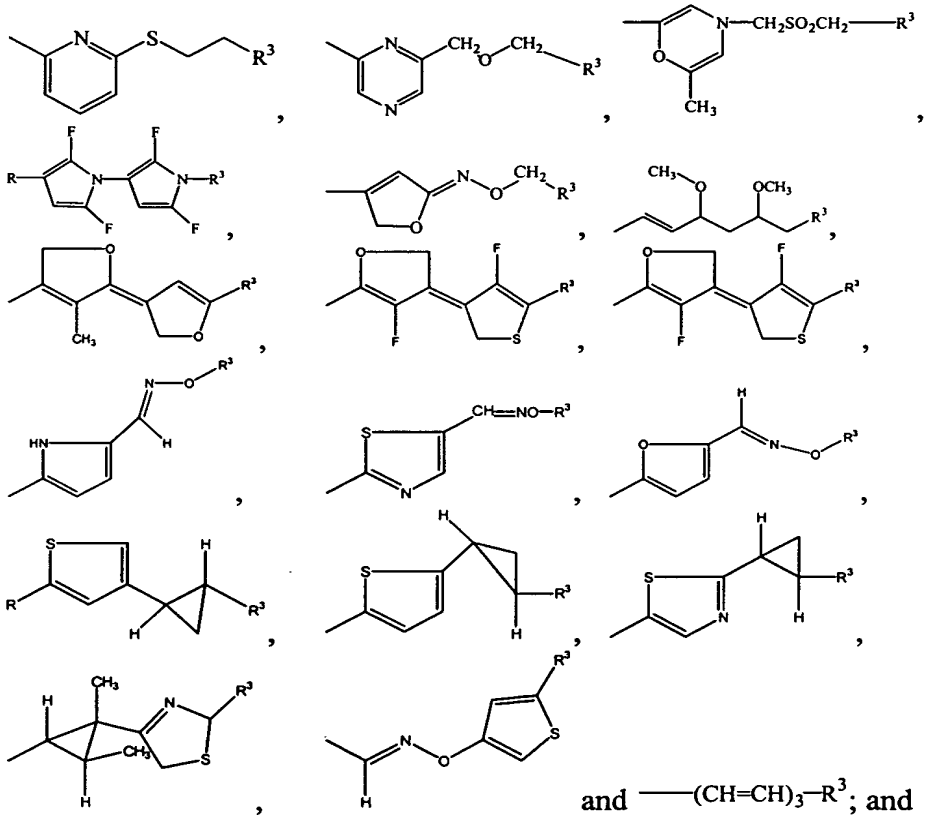
55. The pharmaceutical composition of claim **54**, wherein R^1 is an optionally substituted $(C_2\text{-}C_{18})$alkylene having at least 2 sites of unsaturation.

56. The pharmaceutical composition of claim **54**, wherein said compound is



Ia

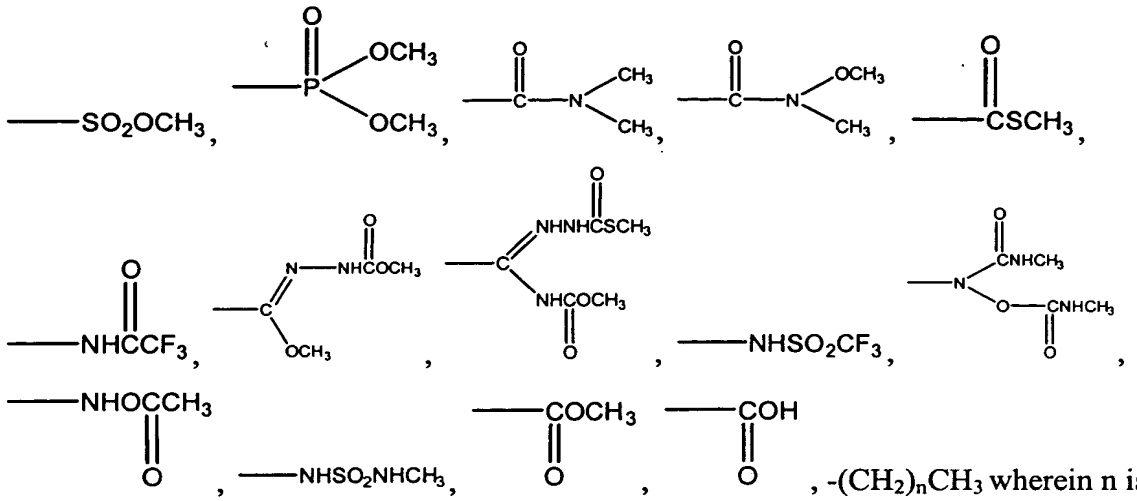
57. The pharmaceutical composition of claim **53**, wherein R^1 is a member selected from the group consisting of



and $-(CH=CH)_3-R^3$; and

R^3 is a member selected from the group consisting of consisting of alkyl, alkoxysulfonyl, dialkylphosphono, optionally substituted carbamoyl, alkylthiocarbonyl, optionally substituted alkylamido, optionally substituted amidino, optionally substituted alkylsulfonamido, alkoyloxyamino, alkylaminosulfonamido, and alkoxycarbonyl.

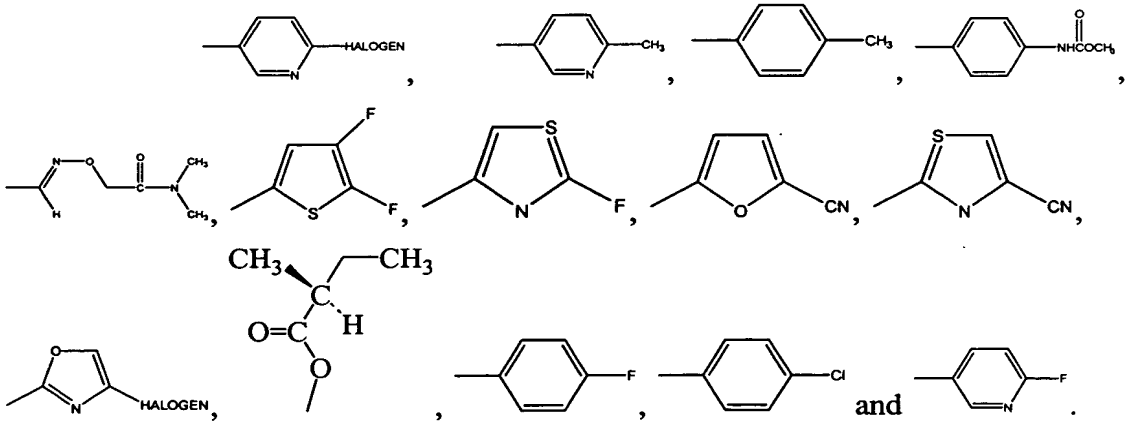
58. The pharmaceutical composition of claim **57**, wherein R^3 is a member selected from the group consisting of consisting of



, $-(CH_2)_nCH_3$ wherein n is about 1 to about 10 and $-CH_3$.

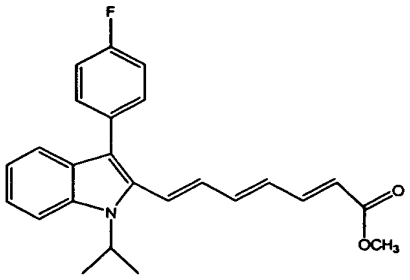
59. The pharmaceutical composition of claim **57**, wherein R^2 is independently a member selected from the group consisting of optionally substituted alkoxy, acyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy.

60. The pharmaceutical composition of claim **57**, wherein R^2 is a member selected from the group consisting of consisting of



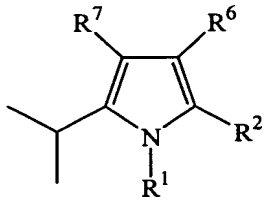
.

61. The pharmaceutical composition of claim **53**, wherein said compound has the formula



62. A pharmaceutical composition, said pharmaceutical composition comprising:

a compound having the formula



II

wherein:

R^1 is a member selected from the group consisting of optionally substituted alkyl, optionally substituted alkoxy, optionally substituted alkylene having at least 2 sites of unsaturation, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, optionally substituted heteroarylthioalkyl, optionally substituted heteroaryliminooxyalkyl, optionally substituted heterocyclyl, optionally substituted oximinoaryl and optionally substituted heteroarylalkoxy;

R^2 is a member selected from the group consisting of optionally substituted $(C_1\text{-}C_6)$alkyl, optionally substituted $(C_1\text{-}C_6)$alkoxy, acyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy;

R^6 is a member selected from the group consisting of an optionally substituted alkyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy;

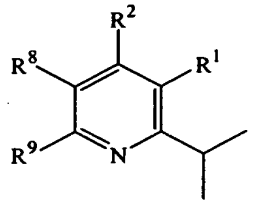
R^7 is a member selected from the group consisting of an optionally substituted alkyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy; or alternatively, R^6 and R^7 and

the carbons to which they are attached, joined to form an optionally substituted aryl or optionally substituted heteroalkyl 5-or 6 membered ring; and

a pharmaceutically acceptable excipient therefor.

63. A pharmaceutical composition, said pharmaceutical composition comprising:

a compound having the formula



III

wherein:

R^1 is a member selected from the group consisting of optionally substituted alkyl, optionally substituted alkoxy, optionally substituted alkylene having at least 2 sites of unsaturation, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, optionally substituted heteroarylthioalkyl, optionally substituted heteroaryliminooxyalkyl, optionally substituted heterocyclyl, optionally substituted oximinoaryl and optionally substituted heteroarylalkoxy;

R^2 is independently a member selected from the group consisting of optionally substituted $(C_1\text{-}C_6)$alkyl, optionally substituted $(C_1\text{-}C_6)$alkoxy, acyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy;

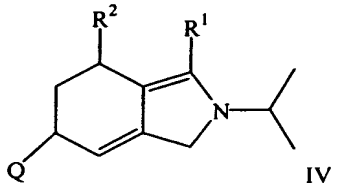
R^8 is a member selected from the group consisting of an optionally substituted alkyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy;

R^9 is a member selected from the group consisting of an optionally substituted alkyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, optionally substituted heteroarylalkoxy; or alternatively, R^8 and R^9 and the carbons to which they are attached, joined to form an optionally substituted aryl or optionally substituted heteroalkyl 5-or 6 membered ring; and

a pharmaceutically acceptable excipient therefor.

64. A pharmaceutical composition, said pharmaceutical composition comprising:

a compound having the formula



IV

wherein:

R^1 is a member selected from the group consisting of optionally substituted alkyl, optionally substituted alkoxy, optionally substituted alkylene having at least 2 sites of unsaturation, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, optionally substituted heteroarylthioalkyl, optionally substituted heteroaryliminooxyalkyl, optionally substituted heterocyclyl, optionally substituted oximinoaryl and optionally substituted heteroarylalkoxy;

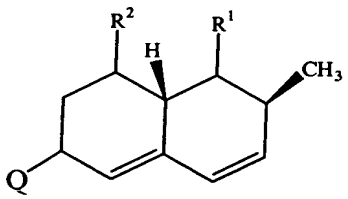
R^2 is a member selected from the group consisting of optionally substituted $(C_1\text{-}C_6)$alkyl, optionally substituted $(C_1\text{-}C_6)$alkoxy, acyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy;

Q is a member selected from the group consisting of hydrogen, optionally substituted alkyl, optionally substituted alkoxy and hydroxy; and

a pharmaceutically acceptable excipient therefor.

65. A pharmaceutical composition, said pharmaceutical composition comprising:

a compound having the formula



V

wherein:

R^1 is a member selected from the group consisting of optionally substituted alkyl, optionally substituted alkoxy, optionally substituted alkylene having at least 2 sites of

unsaturation, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, optionally substituted heteroarylthioalkyl, optionally substituted heteroaryliminooxyalkyl, optionally substituted heterocyclyl, optionally substituted oximinoaryl and optionally substituted heteroarylalkoxy;

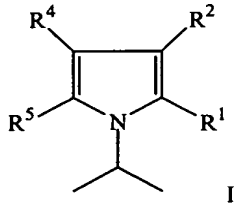
R^2 is independently a member selected from the group consisting of optionally substituted $(C_1\text{-}C_6)$alkyl, optionally substituted $(C_1\text{-}C_6)$alkoxy, acyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy;

Q is a member selected from the group consisting of hydrogen, optionally substituted alkyl, optionally substituted alkoxy and hydroxy; and

pharmaceutically acceptable excipient therefor.

66. A method for blocking calcium influx from the extracellular space, said method comprising: contacting a cell with a store operated calcium influx (SOC) inhibitor, thereby blocking calcium influx from the extracellular space.

67. The method of claim **66**, wherein said SOC inhibitor is a compound having the formula



I

wherein:

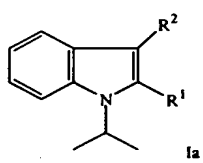
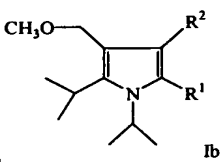
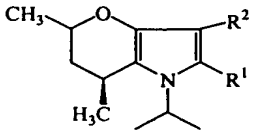
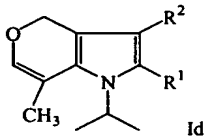
R^1 is a member selected from the group consisting of optionally substituted alkyl, optionally substituted alkoxy, optionally substituted alkylene having at least 2 sites of unsaturation, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, optionally substituted heteroarylthioalkyl, optionally substituted heteroaryliminooxyalkyl, optionally substituted heterocyclyl, optionally substituted oximinoaryl and optionally substituted heteroarylalkoxy;

R^2 is a member selected from the group consisting of optionally substituted $(C_1\text{-}C_6)$alkyl, optionally substituted $(C_1\text{-}C_6)$alkoxy, acyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, optionally substituted heteroarylalkoxy;

R^4 is an optionally substituted alkyl; and

R^5 is an optionally substituted alkyl, or alternatively, R^4 and R^5 and the carbons to which they are attached, joined to form an optionally substituted aryl or optionally substituted heteroalkyl 5-or 6 membered ring.

68. The method of claim **67**, wherein said SOC inhibitor is a compound selected from the group consisting of


Ia,

Ib,

Ic and

Id

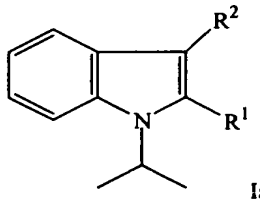
wherein:

R^1 is a member selected from the group consisting of optionally substituted alkyl, optionally substituted alkoxy, optionally substituted (C_2-C_{18})alkylene having at least 2 sites of unsaturation, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, optionally substituted heteroarylalkoxy, and optionally substituted; and

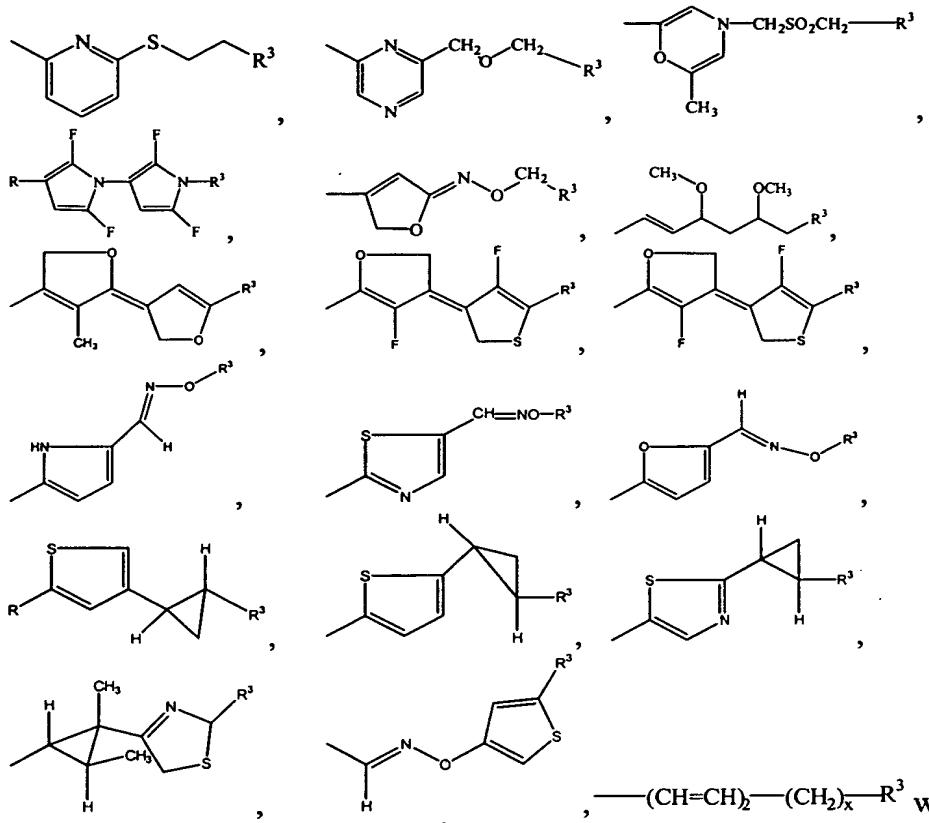
R^2 is independently a member selected from the group consisting of optionally substituted (C_1-C_6)alkyl, optionally substituted (C_1-C_6)alkoxy, acyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy.

69. The method of claim **67**, wherein said SOC inhibitor is a compound wherein R^1 is an optionally substituted (C_2-C_{18})alkylene having at least 2 sites of unsaturation.

70. The method of claim **68**, wherein said compound is


Ia

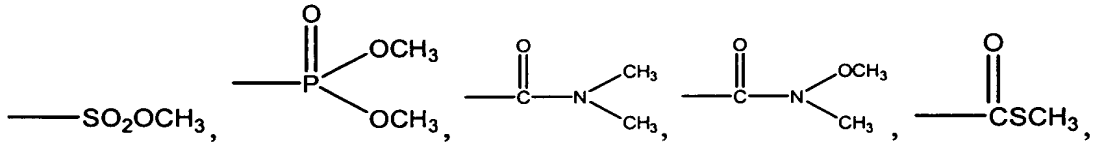
71. The method of claim **67**, wherein R^1 is a member selected from the group consisting of

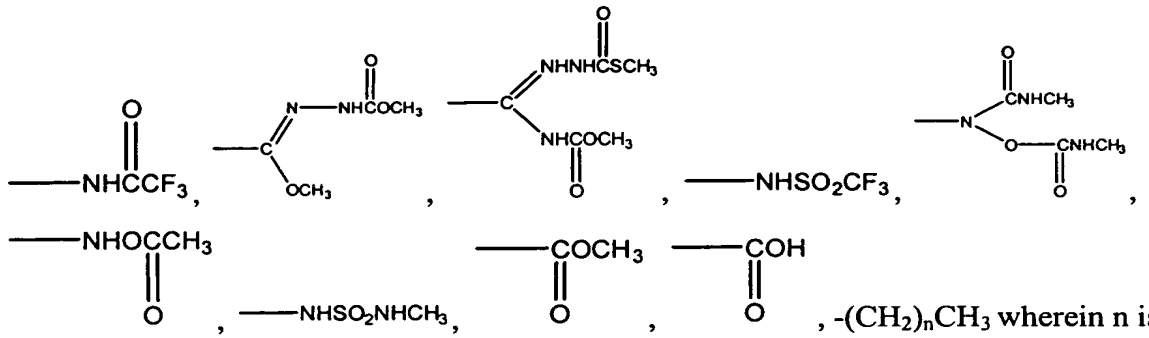


, —(CH=CH)$_2$—(CH$_2$)$_x$—R^3 wherein x is about 1 to about 14 and —(CH=CH)$_3$—R^3; and

R^3 is a member selected from the group consisting of consisting of alkyl, alkoxysulfonyl, dialkylphosphono, optionally substituted carbamoyl, alkylthiocarbonyl, optionally substituted alkylamido, optionally substituted amidino, optionally substituted alkylsulfonamido, alkoyloxyamino, alkylaminosulfonamido, and alkoxycarbonyl.

72. The method of claim **71**, wherein R^3 is a member selected from the group consisting of consisting of

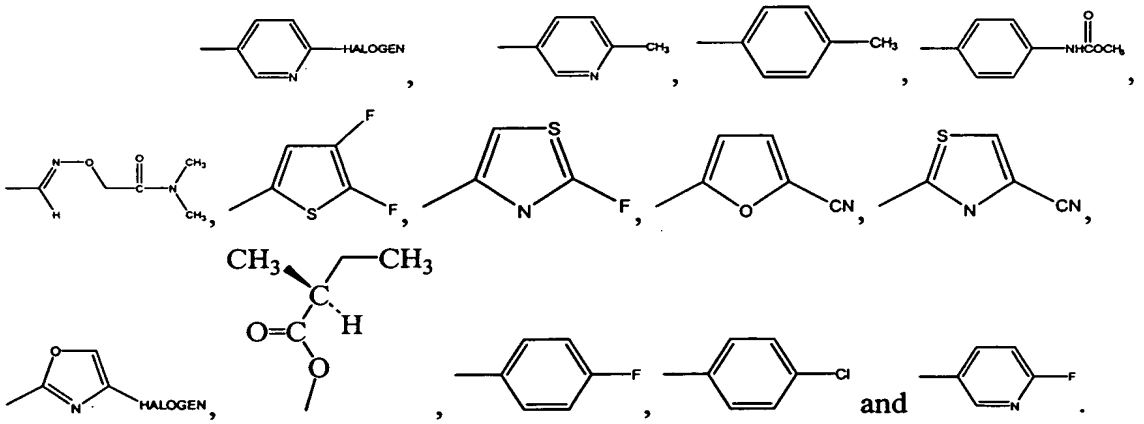




, -$(CH_2)_nCH_3$ wherein n is about 1 to about 10 and -CH_3.

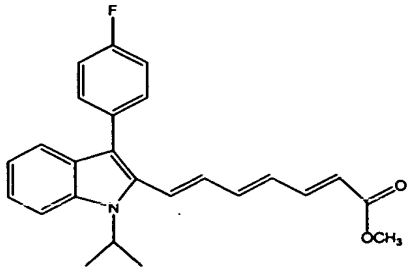
73. The method of claim **71**, wherein R^2 is independently a member selected from the group consisting of optionally substituted alkoxy, acyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy.

74. The method of claim **71**, wherein R^2 is a member selected from the group consisting of consisting of

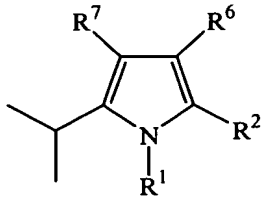


and .

75. The method of claim **67**, wherein said compound has the formula



76. The method of claim 66, wherein said SOC inhibitor is a compound having the formula



II

wherein:

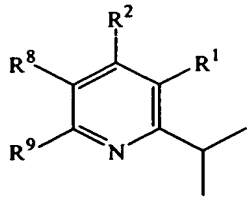
R^1 is a member selected from the group consisting of optionally substituted alkyl, optionally substituted alkoxy, optionally substituted alkylene having at least 2 sites of unsaturation, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, optionally substituted heteroarylthioalkyl, optionally substituted heteroaryliminooxyalkyl, optionally substituted heterocyclyl, optionally substituted oximinoaryl and optionally substituted heteroarylalkoxy;

R^2 is independently a member selected from the group consisting of optionally substituted $(C_1\text{-}C_6)$alkyl, optionally substituted $(C_1\text{-}C_6)$alkoxy, acyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy;

R^6 is a member selected from the group consisting of an optionally substituted alkyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy;

R^7 is a member selected from the group consisting of an optionally substituted alkyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy; or alternatively, R^6 and R^7 and the carbons to which they are attached, joined to form an optionally substituted aryl or optionally substituted heteroalkyl 5-or 6 membered ring.

77. The method of claim 66, wherein said SOC inhibitor is a compound having the formula



III

wherein:

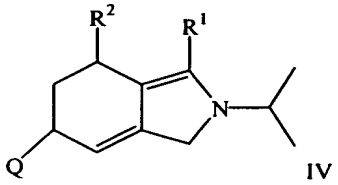
R^1 is a member selected from the group consisting of optionally substituted alkyl, optionally substituted alkoxy, optionally substituted alkylene having at least 2 sites of unsaturation, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, optionally substituted heteroarylthioalkyl, optionally substituted heteroaryliminooxyalkyl, optionally substituted heterocyclyl, optionally substituted oximinoaryl and optionally substituted heteroarylalkoxy;

R^2 is independently a member selected from the group consisting of optionally substituted $(C_1\text{-}C_6)$alkyl, optionally substituted $(C_1\text{-}C_6)$alkoxy, acyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy;

R^8 is a member selected from the group consisting of an optionally substituted alkyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy;

R^9 is a member selected from the group consisting of an optionally substituted alkyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy; or alternatively, R^8 and R^9 and the carbons to which they are attached, joined to form an optionally substituted aryl or optionally substituted heteroalkyl 5-or 6 membered ring.

78. The method of claim **66**, wherein said SOC inhibitor is a compound having the formula



IV

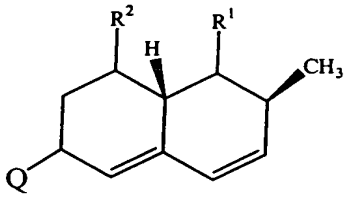
wherein:

R^1 is a member selected from the group consisting of optionally substituted alkyl, optionally substituted alkoxy, optionally substituted alkylene having at least 2 sites of unsaturation, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, optionally substituted heteroarylthioalkyl, optionally substituted heteroaryliminooxyalkyl, optionally substituted heterocyclyl, optionally substituted oximinoaryl and optionally substituted heteroarylalkoxy;

R^2 is a member selected from the group consisting of optionally substituted (C_1-C_6)alkyl, optionally substituted (C_1-C_6)alkoxy, acyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy; and

Q is a member selected from the group consisting of hydrogen, optionally substituted alkyl, optionally substituted alkoxy and hydroxy.

79. The method of claim **66**, wherein said SOC inhibitor is a compound having the formula



V

wherein:

R^1 is a member selected from the group consisting of optionally substituted alkyl, optionally substituted alkoxy, optionally substituted alkylene having at least 2 sites of unsaturation, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, optionally substituted heteroarylthioalkyl, optionally substituted heteroaryliminooxyalkyl, optionally substituted heterocyclyl, optionally substituted oximinoaryl and optionally substituted heteroarylalkoxy;

R^2 is independently a member selected from the group consisting of optionally substituted (C_1-C_6)alkyl, optionally substituted (C_1-C_6)alkoxy, acyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroarylalkyl, and optionally substituted heteroarylalkoxy; and

Q is a member selected from the group consisting of hydrogen, optionally substituted alkyl, optionally substituted alkoxy and hydroxy.

80. The method of claim **66**, wherein said SOC inhibitor is a δ-lactone-containing statin.

81. The method of claim **80**, wherein said δ-lactone-containing statin is a member selected from the group consisting of lovastatin, mevastatin, and simvastatin.

82. A method for treating inflammatory bowel disease (IBD), said method comprising:

administering a store operated calcium influx (SOC) inhibitor, thereby treating inflammatory bowel disease (IBD).

83. A method of treating a disease, comprising administering a pharmaceutical composition comprising an aerosol formulation of a SOC inhibitor, wherein said disease is selected from the group consisting of acute lung injury, adult respiratory distress syndrome, asthma, interstitial lung disease, emphysema, chronic bronchitis and cystic fibrosis.